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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/537,583	12/14/2005	12/14/2005 Katherine Ann Vousden		4893
23565 KLAUBER &	7590 12/27/2007		EXAMINER	
411 HACKEN	SACK AVENUE		OGUNBIYI, OLUWATOSIN A	
HACKENSACK, NJ 07601			ART UNIT	PAPER NUMBER
			1645	
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)		
	10/537,583	VOUSDEN, KATHERINE ANN		
Office Action Summary	Examiner	Art Unit		
	Oluwatosin Ogunbiyi	1645		
The MAILING DATE of this communication	appears on the cover sheet wit	h the correspondence address		
Period for Reply	DIVIO OFT TO EVENE A MAC	ONTHE OF THE TY (OO) DAYS		
A SHORTENED STATUTORY PERIOD FOR RE WHICHEVER IS LONGER, FROM THE MAILING - Extensions of time may be available under the provisions of 37 CFI after SIX (6) MONTHS from the mailing date of this communication - If NO period for reply is specified above, the maximum statutory pe - Failure to reply within the set or extended period for reply will, by st Any reply received by the Office later than three months after the mearned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNIC R 1.136(a). In no event, however, may a re riod will apply and will expire SIX (6) MONT atute, cause the application to become ABA	ATION. ply be timely filed "HS from the mailing date of this communication. ANDONED (35 U.S.C. § 133).		
Status				
1)⊠ Responsive to communication(s) filed on <u>0</u>	<u> 5 November 2007</u> .			
2a) ☐ This action is FINAL . 2b) ☑ 7	This action is FINAL . 2b)⊠ This action is non-final.			
3) Since this application is in condition for allo	wance except for formal matte	ers, prosecution as to the merits is		
closed in accordance with the practice und	er <i>Ex parte Quayle</i> , 1935 C.D.	11, 453 O.G. 213.		
Disposition of Claims				
4)⊠ Claim(s) <u>1,8,14-16 and 18-22</u> is/are pendin	g in the application.			
4a) Of the above claim(s) 14-16 and 18-22		ration.		
5) Claim(s) is/are allowed.				
6)⊠ Claim(s) <u>1 and 8</u> is/are rejected.				
7) Claim(s) <u>8</u> is/are objected to.				
8) Claim(s) are subject to restriction ar	id/or election requirement.			
Application Papers				
9)☐ The specification is objected to by the Exam	niner.			
10)☐ The drawing(s) filed on is/are: a)☐ :	accepted or b)⊡ objected to b	y the Examiner.		
Applicant may not request that any objection to	= · · ·			
Replacement drawing sheet(s) including the co				
11)☐ The oath or declaration is objected to by the	e Examiner, Note the attached	Office Action of form PTO-192.		
Priority under 35 U.S.C. § 119				
12) Acknowledgment is made of a claim for fore	eign priority under 35 U.S.C. §	119(a)-(d) or (f).		
a) ☐ All b) ☐ Some * c) ☐ None of:				
1. Certified copies of the priority docum				
2. Certified copies of the priority docum	·			
 Copies of the certified copies of the papelication from the International But 	•	eceived in this National Stage		
* See the attached detailed Office action for a		received		
2222				
Attachment(s)	_			
 Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-948) 		ummary (PTO-413))/Mail Date		
 2) Interpretation Disclosure Statement(s) (PTO/SB/08) 	5) 🔲 Notice of Int	formal Patent Application		
Paper No(s)/Mail Date	6)	<u> -</u> ·		

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DETAILED ACTION

Claims 1, 8, 14-16 and 18-22 are pending. Claims 2-7, 9-10 and 17 are canceled.

Claims 1 and 8 are under examination

REQUEST FOR CONTINUED EXAMINATION

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 11/5/07 has been entered.

Claim Objections

Claim 8 is objected to because of the following informalities: 'albicans' is misspelled.

Appropriate correction is required.

Claim Rejections - 35 USC § 112

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly

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> connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claim 8 is rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the enablement requirement. The claim(s) contains subject matter, which was not described in the specification in such a way as to enable one skilled in the art to which it pertains, or with which it is most nearly connected, to make and/or use the invention.

The claim is drawn a method of screening or testing for candidate anti-fungal compounds that impair *Candida albicans* ATP (CTP): tRNA nucleotidyltransferase enzyme (CCA1) activity comprising:

- a) providing a *C. albicans* cell wherein the cell expresses *Candida albicans*ATP (CTP): tRNA nucleotidyltransferase enzyme (CCA1) under the control of a heterologous promoter
- (b) providing one or more candidate compounds;
- c) contacting said *Candida albicans* cell(s) with said one or more candidate compounds;
- d) inducing said promoter;
- e) determining whether the candidate compound is a CCA1 inhibitor.

The nature of the invention as set forth supra involves screening or testing for an antifungal compound that impairs *C. albicans* CCA1 in a *Candida albicans* cell expressing CCA1 under the control of a heterologous promoter by determining whether the candidate compound is a CCA 1 inhibitor.

The specification on p.2 line 38 teaches that CCA1 is an essential protein for the fungal species *Candida*. Example 4 on p. 10 provides a demonstration of CCA1 as an essential gene

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product in Candida albicans. A Candida albicans strain with one copy of the CCA1 gene under the control of tetracycline inducible promoter expresses CCA1 under strong induction i.e. without tetracycline and does not express CCA1 under tight repression i.e. with tetracycline. Under CCA1 expression, there is cell growth and no significant growth was observed when CCA1 is repressed (p. 11 example 4.2).

The method as claimed does not provide for how to determine whether the candidate compound is a CCA1 inhibitor after contacting said C. albicans cell expressing C. albicans CCA1 with said one or more candidate compounds.

For this claimed method the specification only gives guidance as to how to determine that a candidate compound is a CCA1 inhibitor in an eukaryotic cell (such as C. albicans) expressing CCA1 (p. 5 lines 23-30). The method entails that after contacting said C. albicans cell expressing C. albicans CCA1 with said one or more candidate compounds, the interaction of the candidate compound with CCA1 is assessed by determining the effect on growth and viability of said cells.

However, this in vivo assay system for screening or testing for an anti-fungal compound that impairs C. albicans CCA1 function does not take into account that there are more than one essential genes in C. albicans (Song et al. Microbiology (2003), 149:219-259, Veses et al. Eukaryotic Cell (2005), p. 1088-1101, Bruno et al. Trends in Microbiology (2004) 12: 157-161, p.159, table 2) and also a compound may have multiple targets including CCA1. The claim requires contacting a candidate compound with a Candida albicans cell(s) expressing CCA1. If negative effects on growth or viability are seen in said cell(s), how does one of skill in the art determine that said negative effect is due to impairment of CCA1 by the candidate compound? The specification does not correlate impairment of growth or viability with the direct impairment

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with CCA1. The assay described in the specification does not provide any guidance or direction as to how to rule out the effects of such compound on other essential genes in said *C. albicans* expressing CCA1. Further, this in vivo assay for screening or testing for an anti-fungal compound that impairs *C. albicans* CCA1 and that described in the claim does not take into account that there is endogenous CCA1 in the cells expressing *Candida albicans* CCA1.

For example, Onishi et al (Feb. 2000, Antimicrobial Agents and Chemotherapy p. 368-377 cited in previous action) teaches a screen for in vitro antifungal activity of several compounds by a growth inhibition assay (page 369 column 1 materials and methods and table 1) and then the compounds were evaluated to determine whether said compounds were direct inhibitors of the enzyme by measuring the enzyme's activity in the presence of said compounds (page 370 column 2 first full paragraph, page 373 column 1 - 2 and table 4). Onishi et al takes into account that the fungal cell expresses the enzyme naturally and Onishi et al do not over express the enzyme in the same fungal cell. Instead Onishi et al assesses growth inhibition and then screens for enzyme activity in vitro in the presence of the potential anti-fungal.

Assessing inhibition of growth does not provide any knowledge about the effect of the compound on CCA1 activity because as mentioned above a compound may have more that one target in said *C. albicans* expressing CCA1 including CCA1. Furthermore, an anti-fungal compound has many different activities (see Ghannoum et al. 1999. Clinical Microbiology Reviews, p. 501-517 for different mode of actions of some anti-fungal compounds, cited in previous office action) and these compounds would inhibit growth but do not have to impair activity of CCA1. As such, impairment of growth is not directly correlated with impairment of CCA1 activity in the method of screening set forth in the specification i.e. method of determining whether the candidate compound inhibits CCA1 in the C. albicans cells expressing CCA1. Claim 8 does not teach how to determine whether the candidate compound is a CCA1 inhibitor.

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The method for performing such determination as disclosed in the specification and set forth above i.e. assessing effects on growth and viability does not does not correlate impairment of growth or viability with the direct impairment of CCA1 function or activity.

In view of the above, it would require undue experimentation for the skilled artisan to use the invention as claimed.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(e) the invention was described in (1) an application for patent, published under section 122(b), by another filed in the United States before the invention by the applicant for patent or (2) a patent granted on an application for patent by another filed in the United States before the invention by the applicant for patent, except that an international application filed under the treaty defined in section 351(a) shall have the effects for purposes of this subsection of an application filed in the United States only if the international application designated the United States and was published under Article 21(2) of such treaty in the English language.

Claim 1 is rejected under 35 U.S.C. 102(e) as being anticipated by Weinstock et al. US 6,747,137 B1 published Jun. 8, 2004, filed Feb 12, 1999.

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The claims are drawn to a method of screening or testing for candidate anti-fungal compounds that impair *Candida albicans ATP(CTP):tRNA* nucleotidyltransferase enzyme (CCA1) activity

comprising:

- a) providing fungal Candida albicans CCA 1;
- b) providing one or more candidate compounds;
- c) contacting said CCA 1 with said one or more candidate compounds; and
- d) determining the ability of the candidate compound to inhibit CCA1 activity.

Weinstock et al teaches a method of screening test compounds for anti-fungal activity comprising providing a *Candida albicans* target sequence such as *Candida albicans* tRNA nucleotidyl transferase also known as CCA1 (table 2 columns 587 and 588 contig3807) and contacting a test compound and determining binding of the test compound to said CCA1 to determine whether said compound has anti-fungal activity (i.e. whether anti-fungal inhibits CCA1 activity). See column 10 lines 28-45, column 20 lines 46-67 to column 21 lines 1-54 (for description of table 2 which discloses *Candida albicans* CCA1).

Status of Claims

Claims 1 and 8 are rejected.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Oluwatosin Ogunbiyi whose telephone number is 571-272-9939. The examiner can normally be reached on M-F 7am-4pm.

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If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Shanon Foley can be reached on 571-272-0898. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Tosin Osmayi

PATRICIA A DUFFY PRIMARY EXAMINER